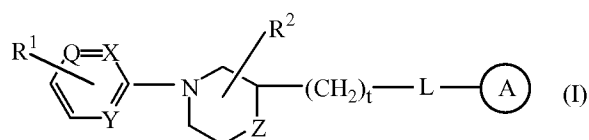


Listing of Claims

This listing of claims will replace all prior versions, and listings, of claims in the application:

Claims 1-27 (cancelled).

28. (New) A compound of formula (I),



the pharmaceutically acceptable addition salts and the stereo-chemically isomeric forms thereof, wherein

t is 0, 1, 2, 3 or 4 and when t is 0 then a direct bond is intended;

each Q is $\text{—C}\begin{smallmatrix} \diagup \\ \diagdown \end{smallmatrix}$;

each X is nitrogen;

each Y is nitrogen;

each Z is —NH-, or —O-;

R¹ is —C(O)NR³R⁴, —NHC(O)R⁷, —C(O)—C₁₋₆alkanediylSR⁷, —NR⁸C(O)N(OH)R⁷, —NR⁸C(O)C₁₋₆alkanediylSR⁷, or —NR⁸C(O)C=N(OH)R⁷ wherein R³ and R⁴ are each independently selected from hydrogen, hydroxy, C₁₋₆alkyl, hydroxyC₁₋₆alkyl, aminoC₁₋₆alkyl or aminoaryl;

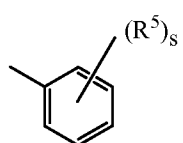
R^7 is independently selected from hydrogen, C_{1-6} alkyl, or C_{1-6} alkylcarbonyl;

R^8 is independently selected from hydrogen or C_{1-6} alkyl;

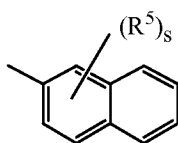
R^2 is hydrogen, hydroxy, amino, hydroxy C_{1-6} alkyl, C_{1-6} alkyl, C_{1-6} alkyloxy, aryl C_{1-6} alkyl, aminocarbonyl, hydroxycarbonyl, amino C_{1-6} alkyl, aminocarbonyl C_{1-6} alkyl, hydroxycarbonyl C_{1-6} alkyl, hydroxyaminocarbonyl, C_{1-6} alkyloxycarbonyl, C_{1-6} alkylamino C_{1-6} alkyl or di(C_{1-6} alkyl)amino C_{1-6} alkyl;

-L- is a bivalent radical selected from $-NR^9C(O)-$, $-NR^9SO_2-$ or $-NR^9CH_2-$ wherein R^9 is hydrogen, C_{1-6} alkyl, C_{3-10} cycloalkyl, hydroxy C_{1-6} alkyl, C_{1-6} alkyloxy C_{1-6} alkyl or di(C_{1-6} alkyl)amino C_{1-6} alkyl;

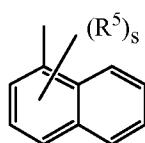
— is a radical selected from



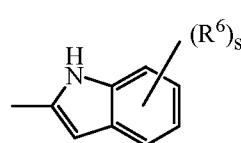
(a-1)



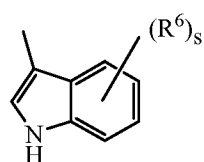
(a-20)



(a-21)

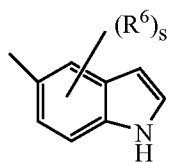


(a-28)



(a-29)

or



(a-48)

;

wherein each s is independently 0, 1, 2, 3, 4 or 5;

R^5 is selected from hydrogen or phenyl optionally substituted with one, two or three substituents independently selected from halo, amino, nitro, C_{1-6} alkyl, C_{1-6} alkyloxy,

hydroxyC₁₋₄alkyl, trifluoromethyl, trifluoromethyloxy, hydroxyC₁₋₄alkyloxy, C₁₋₄alkylsulfonyl, C₁₋₄alkyloxyC₁₋₄alkyloxy, C₁₋₄alkyloxycarbonyl, aminoC₁₋₄alkyloxy, di(C₁₋₄alkyl)aminoC₁₋₄alkyloxy, di(C₁₋₄alkyl)amino, di(C₁₋₄alkyl)aminocarbonyl, di(C₁₋₄alkyl)aminoC₁₋₄alkyl, di(C₁₋₄alkyl)aminoC₁₋₄alkylaminoC₁₋₄alkyl, di(C₁₋₄alkyl)amino(C₁₋₄alkyl)amino, di(C₁₋₄alkyl)amino(C₁₋₄alkyl)aminoC₁₋₄alkyl, di(C₁₋₄alkyl)aminoC₁₋₄alkyl(C₁₋₄alkyl)amino, di(C₁₋₄alkyl)aminoC₁₋₄alkyl(C₁₋₄alkyl)aminoC₁₋₄alkyl, aminosulfonylamino(C₁₋₄alkyl)amino, aminosulfonylamino(C₁₋₄alkyl)aminoC₁₋₄alkyl, di(C₁₋₄alkyl)aminosulfonylamino(C₁₋₄alkyl)amino, di(C₁₋₄alkyl)aminosulfonylamino(C₁₋₄alkyl)aminoC₁₋₆alkyl, cyano, (hydroxyC₁₋₄alkyl)(C₁₋₄alkyl)amino, (hydroxyC₁₋₄alkyl)(C₁₋₄alkyl)aminoC₁₋₄alkyl, hydroxyC₁₋₄alkylaminoC₁₋₄alkyl, di(hydroxyC₁₋₄alkyl)aminoC₁₋₄alkyl, hydroxyC₁₋₄alkylamino, di(hydroxyC₁₋₄alkyl)amino, or di(C₁₋₄alkyl)aminoC₁₋₄alkylamino;


R⁶ is selected from hydrogen; halo; hydroxy; amino; nitro; trihaloC₁₋₆alkyl; trihaloC₁₋₆alkyloxy; C₁₋₆alkyl; C₁₋₆alkyl substituted with aryl and C₃₋₁₀cycloalkyl; C₁₋₆alkyloxy; C₁₋₆alkyloxyC₁₋₆alkyloxy; C₁₋₆alkylcarbonyl; C₁₋₆alkyloxycarbonyl; C₁₋₆alkylsulfonyl; cyanoC₁₋₆alkyl; hydroxyC₁₋₆alkyl; hydroxyC₁₋₆alkyloxy; hydroxyC₁₋₆alkylamino; aminoC₁₋₆alkyloxy; di(C₁₋₆alkyl)aminocarbonyl; di(hydroxyC₁₋₆alkyl)amino; (aryl)(C₁₋₆alkyl)amino; di(C₁₋₆alkyl)aminoC₁₋₆alkyloxy; di(C₁₋₆alkyl)aminoC₁₋₆alkylamino; di(C₁₋₆alkyl)aminoC₁₋₆alkylaminoC₁₋₆alkyl; arylsulfonyl; arylsulfonylamino; aryloxy; aryloxyC₁₋₆alkyl; arylC₂₋₆alkenediyl; di(C₁₋₆alkyl)amino; di(C₁₋₆alkyl)aminoC₁₋₆alkyl; di(C₁₋₆alkyl)amino(C₁₋₆alkyl)amino; di(C₁₋₆alkyl)amino(C₁₋₆alkyl)aminoC₁₋₆alkyl; di(C₁₋₆alkyl)aminoC₁₋₆alkyl(C₁₋₆alkyl)amino; di(C₁₋₆alkyl)aminoC₁₋₆alkyl(C₁₋₆alkyl)aminoC₁₋₆alkyl; aminosulfonylamino(C₁₋₆alkyl)amino; aminosulfonylamino(C₁₋₆alkyl)aminoC₁₋₆alkyl;

di(C₁₋₆alkyl)aminosulfonylamino(C₁₋₆alkyl)amino;
 di(C₁₋₆alkyl)aminosulfonylamino(C₁₋₆alkyl)aminoC₁₋₆alkyl; cyano;
 (hydroxyC₁₋₆alkyl)(C₁₋₆alkyl)amino; (hydroxyC₁₋₆alkyl)(C₁₋₆alkyl)aminoC₁₋₆alkyl;
 hydroxyC₁₋₆alkylaminoC₁₋₆alkyl; di(hydroxyC₁₋₆alkyl)aminoC₁₋₆alkyl;
 phenyl; phenyl substituted with one, two or three substituents independently
 selected from halo, amino, nitro, C₁₋₆alkyl, C₁₋₆alkyloxy,
 hydroxyC₁₋₄alkyl, trifluoromethyl, trifluoromethyloxy, hydroxyC₁₋₄alkyloxy,
 C₁₋₄alkylsulfonyl, C₁₋₄alkyloxyC₁₋₄alkyloxy, C₁₋₄alkyloxycarbonyl,
 aminoC₁₋₄alkyloxy,
 di(C₁₋₄alkyl)aminoC₁₋₄alkyloxy, di(C₁₋₄alkyl)amino, di(C₁₋₄alkyl)aminocarbonyl,
 di(C₁₋₄alkyl)aminoC₁₋₄alkyl, di(C₁₋₄alkyl)aminoC₁₋₄alkylaminoC₁₋₄alkyl,
 di(C₁₋₄alkyl)amino(C₁₋₄alkyl)amino, di(C₁₋₄alkyl)amino(C₁₋₄alkyl)aminoC₁₋₄alkyl,
 di(C₁₋₄alkyl)aminoC₁₋₄alkyl(C₁₋₄alkyl)amino,
 di(C₁₋₄alkyl)aminoC₁₋₄alkyl(C₁₋₄alkyl)aminoC₁₋₄alkyl,
 aminosulfonylamino(C₁₋₄alkyl)amino,
 aminosulfonylamino(C₁₋₄alkyl)aminoC₁₋₄alkyl,
 di(C₁₋₄alkyl)aminosulfonylamino(C₁₋₄alkyl)amino,
 di(C₁₋₄alkyl)aminosulfonylamino(C₁₋₄alkyl)aminoC₁₋₆alkyl, cyano,
 (hydroxyC₁₋₄alkyl)(C₁₋₄alkyl)amino, (hydroxyC₁₋₄alkyl)(C₁₋₄alkyl)aminoC₁₋₄alkyl,
 hydroxyC₁₋₄alkylaminoC₁₋₄alkyl, di(hydroxyC₁₋₄alkyl)aminoC₁₋₄alkyl,
 hydroxyC₁₋₄alkylamino, di(hydroxyC₁₋₄alkyl)amino, or
 di(C₁₋₄alkyl)aminoC₁₋₄alkylamino;

aryl in the above is phenyl, or phenyl substituted with one or more substituents each
 independently selected from halo, C₁₋₆alkyl, C₁₋₆alkyloxy, trifluoromethyl, cyano or
 hydroxycarbonyl.

29. (New) A compound as claimed in claim 28 wherein

R³ and R⁴ are each independently selected from hydrogen, hydroxy, hydroxyC₁₋₆alkyl, aminoC₁₋₆alkyl or aminoaryl;

— is a radical selected from (a-1), (a-20), (a-21), (a-28), or (a-29);

R⁵ is selected from hydrogen; phenyl; or phenyl substituted with one, two or three substituents independently selected from halo, amino, C₁₋₆alkyl, C₁₋₆alkyloxy, hydroxyC₁₋₄alkyl, trifluoromethyl, trifluoromethyloxy, hydroxyC₁₋₄alkyloxy, C₁₋₄alkyloxyC₁₋₄alkyloxy, aminoC₁₋₄alkyloxy, di(C₁₋₄alkyl)aminoC₁₋₄alkyloxy, di(C₁₋₄alkyl)amino, di(C₁₋₄alkyl)aminoC₁₋₄alkyl, di(C₁₋₄alkyl)aminoC₁₋₄alkyl(C₁₋₄alkyl)aminoC₁₋₄alkyl, hydroxyC₁₋₄alkylamino, di(hydroxyC₁₋₄alkyl)amino, or di(C₁₋₄alkyl)aminoC₁₋₄alkylamino.

R⁶ is selected from hydrogen; halo; hydroxy; amino; nitro; trihaloC₁₋₆alkyl; trihaloC₁₋₆alkyloxy; C₁₋₆alkyl; C₁₋₆alkyloxy; C₁₋₆alkyloxyC₁₋₆alkyloxy; C₁₋₆alkylcarbonyl; C₁₋₆alkylsulfonyl; cyanoC₁₋₆alkyl; hydroxyC₁₋₆alkyl; hydroxyC₁₋₆alkyloxy; hydroxyC₁₋₆alkylamino; aminoC₁₋₆alkyloxy; di(C₁₋₆alkyl)aminocarbonyl; di(hydroxyC₁₋₆alkyl)amino; arylC₁₋₆alkyl)amino; di(C₁₋₆alkyl)aminoC₁₋₆alkyloxy; di(C₁₋₆alkyl)aminoC₁₋₆alkylamino; arylsulfonyl; arylsulfonylamino; aryloxy; arylC₂₋₆alkenediyl; di(C₁₋₆alkyl)amino; di(C₁₋₆alkyl)aminoC₁₋₆alkyl; di(C₁₋₆alkyl)aminoC₁₋₆alkyl(C₁₋₆alkyl)aminoC₁₋₆alkyl; cyano; thiophenyl; thiophenyl substituted with di(C₁₋₆alkyl)aminoC₁₋₆alkyl(C₁₋₆alkyl)aminoC₁₋₆alkyl, di(C₁₋₆alkyl)aminoC₁₋₆alkyl, or di(hydroxyC₁₋₆alkyl)aminoC₁₋₆alkyl; (hydroxyC₁₋₆alkyl)(C₁₋₆alkyl)amino; (hydroxyC₁₋₆alkyl)(C₁₋₆alkyl)aminoC₁₋₆alkyl; or phenyl optionally substituted with one, two or three substituents independently selected from halo, amino, C₁₋₆alkyl, C₁₋₆alkyloxy, hydroxyC₁₋₄alkyl, trifluoromethyl, trifluoromethyloxy, hydroxyC₁₋₄alkyloxy, C₁₋₄alkyloxyC₁₋₄alkyloxy, aminoC₁₋₄alkyloxy, di(C₁₋₄alkyl)aminoC₁₋₄alkyloxy, di(C₁₋₄alkyl)amino, di(C₁₋₄alkyl)aminoC₁₋₄alkyl, di(C₁₋₄alkyl)aminoC₁₋₄alkyl(C₁₋₄alkyl)aminoC₁₋₄alkyl, (hydroxyC₁₋₄alkyl)(C₁₋₄alkyl)amino, (hydroxyC₁₋₄alkyl)(C₁₋₄alkyl)aminoC₁₋₄alkyl, hydroxyC₁₋₄alkylamino, di(hydroxyC₁₋₄alkyl)amino, or di(C₁₋₄alkyl)aminoC₁₋₄alkylamino.

30. (New) A compound as claimed in claim 28 wherein t is 0;

R^1 is $-C(O)NR^3R^4$, $-C(O)-C_{1-6}alkanediyISR^7$, $-NR^8C(O)N(OH)R^7$,
 $-NR^8C(O)C_{1-6}alkanediyISR^7$ or $-NR^8C(O)C=N(OH)R^7$ wherein R^3 and R^4 are each
independently selected from hydrogen, hydroxy, hydroxy $C_{1-6}alkyl$ or amino $C_{1-6}alkyl$;
 R^2 is hydrogen, hydroxy, amino,
hydroxy $C_{1-6}alkyl$, $C_{1-6}alkyl$, $C_{1-6}alkyloxy$, aryl $C_{1-6}alkyl$, aminocarbonyl,
amino $C_{1-6}alkyl$, $C_{1-6}alkylaminoC_{1-6}alkyl$ or di($C_{1-6}alkyl$)amino $C_{1-6}alkyl$;
-L- is a bivalent radical selected from $-NHC(O)-$ or $-NHSO_2-$;

$\text{---}\textcircled{A}$ is a radical selected from (a-1), (a-20),
(a-21), (a-28), or (a-48);

each s is independently 0, 1, 2, 3 or 4;

R^5 is hydrogen, phenyl or phenyl substituted with one or two substituents
independently selected from halo, $C_{1-6}alkyl$, $C_{1-6}alkyloxy$ or trifluoromethyl;
and R^6 is hydrogen, phenyl; or phenyl substituted with one or two substituents
independently selected from halo, $C_{1-6}alkyl$, $C_{1-6}alkyloxy$ or trifluoromethyl.

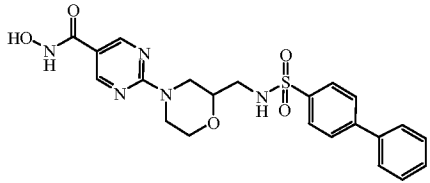
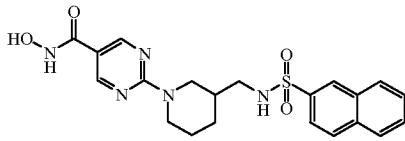
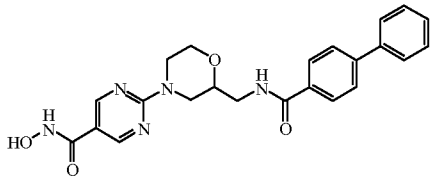
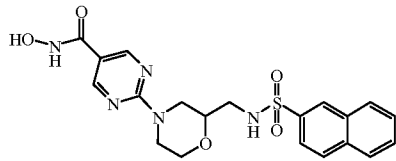
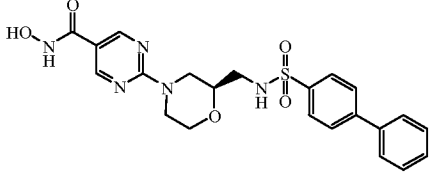
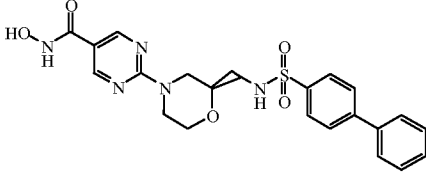
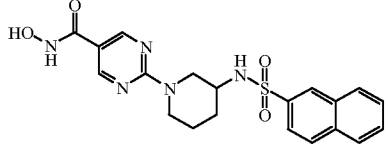
31. (New) A compound as claimed in claim 28 wherein t is 0 or 1; each Q is $\text{---}\text{C}\text{=}\text{C}\text{=}$;
each X is nitrogen; R^1 is $-C(O)NH(OH)$; R^2 is hydrogen, hydroxy, $C_{1-6}alkyl$, or
aryl $C_{1-6}alkyl$; -L- is a bivalent radical selected from $-NHC(O)-$ or $-NHSO_2-$;

$\text{---}\textcircled{A}$ is a radical selected from (a-1) or (a-20); each s is independently 0 or 1;
and each R^5 is independently selected from hydrogen or phenyl.

32. (New) A compound as claimed in claim 28 wherein t is 1; each Q is $\text{---}\text{C}\text{=}\text{C}\text{=}$; each
X is nitrogen; each Y is nitrogen; each Z is $-O-$; R^1 is $-C(O)NH(OH)$; R^2 is

hydrogen; -L- is a bivalent radical selected from $-NHC(O)-$ or $-NHSO_2-$; $\text{---}\textcircled{A}$
is a radical selected from (a-1) or (a-20); each s is independently 0 or 1; and each
 R^5 is independently selected from hydrogen or phenyl.

33.(New) A compound according to claim 28 selected from the following compounds No 4, No 10, No 8, No 6, No 1, No 12 and No 14 :

 <p>Co. No. 4</p>	 <p>Co. No. 10</p>
 <p>Co. No. 8</p>	 <p>Co. No. 6</p>
 <p>Co. No. 1</p>	 <p>Co. No. 12</p>
 <p>Co. No. 14.</p>	

34. (New) A pharmaceutical composition comprising pharmaceutically acceptable carriers and as an active ingredient a therapeutically effective amount of a compound as claimed in claim 28.

35. (New) A combination of anti-cancer agents and a compound of claim 28.

36. (New) A pharmaceutical composition comprising pharmaceutically acceptable carriers and as an active ingredient a therapeutically effective amount of a compound as claimed in claim 33.

37. (New) A combination of anti-cancer agents and a compound of claim 33.